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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/541,875	04/05/2006	Jurgen Dorn	14673-070US	7921
79990	7590	10/01/2009		
C. R. Bard, Inc. Bard Peripheral Vascular, Inc. 1415 W. 3rd Street P.O. Box 1740 Tempe, AZ 85280-1740			EXAMINER HORNBERGER, JENNIFER LEA	
			ART UNIT 3734	PAPER NUMBER
			MAIL DATE 10/01/2009	DELIVERY MODE PAPER

**Please find below and/or attached an Office communication concerning this application or proceeding.**

The time period for reply, if any, is set in the attached communication.



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## **DETAILED ACTION**

### ***Election/Restrictions***

1. This application contains claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

The species are as follows:

- I. Surgical delivery device as disclosed with respect to Figures 1, 2A, and 2B.
- II. Surgical delivery device as disclosed with respect to Figures 3, 3A, 3B, and 4.
- III. Surgical delivery device as disclosed with respect to Figures 5 and 6.

Applicant is required, in reply to this action, to elect a single species to which the claims shall be restricted if no generic claim is finally held to be allowable. The reply must also identify the claims readable on the elected species, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered non-responsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

2. The claims are deemed to correspond to the species listed above in the following manner:

- Species I. Claims 1-26 and 39-40.
- Species II. Claims 1, 2, 7, 8, 10, 11, and 27-38.
- Species III. Claims 1-8, 10, and 11.

The following claim(s) are generic: 1, 2, 7, 8, 10, and 11.

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3. The species listed above do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, the species lack the same or corresponding special technical features for the following reasons: The common technical feature is a delivery device having a proximal end, a primary shaft which is a tube, and a distal zone to be advanced over a guidewire along a bodily lumen to a site of surgery, the device characterized by a tubular means for defining a guidewire lumen, said tubular means lying within the distal zone with the guidewire lumen to one side of the primary shaft and having a proximal end opening which lies to one side of the shaft; a sleeve shaped means for defining a lumen to receive a surgical element distal of the tubular means, the sleeve shaped means having a proximal end which is form-fitted over the primary shaft and has a radially inwardly tapering portion proximal of the proximal end of the tubular means, said inwardly tapering portion defining a proximal guidewire lumen exit port.

Keegan et al. (US 2003/0109886) disclose a delivery device (Figures 17-19(d)) having a proximal end, a tubular primary shaft (2), and a distal zone to be advanced over a guidewire along a bodily lumen to a site of surgery, the device characterized by a tubular means for defining a guidewire lumen (101), said tubular means lying within the distal zone with the guidewire lumen to one side of the primary shaft and having a proximal opening which lies to one side of the shaft, a sleeve shaped means (4) for defining a lumen to receive a surgical element distal of the tubular means, the sleeve shaped means having a braided reinforcing filamentary material in the distal end (paragraph 119) and having a proximal end which is form-fitted over the primary shaft and has a radially inwardly tapering portion (9) proximal of the proximal end of the tubular means, said inwardly tapering portion defining a proximal guidewire lumen exit port (11).

4. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to JENNIFER L. HORNBERGER whose telephone number is (571)270-3642. The examiner can normally be reached on Monday through Friday from 8am-5pm, Eastern Time.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Todd Manahan can be reached on (571)272-4713. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/J. L. H./  
Examiner, Art Unit 3734

/Todd E Manahan/  
Supervisory Patent Examiner, Art Unit 3734